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(71) Applicant (for all designated States except US): **ARI-
ZONA BOARD OF REGENTS**, acting for and on
behalf of, **ARIZONA STATE UNIVERSITY** [US/US];
c/o Arizona Technology Enterprises LLC, Brickyard Suite
601, Room 691AA, 699 S. Mill Avenue, Tempe, AZ 85281
(US).

(71) Applicant and

(72) Inventor: **FRASCH, Wayne, D.** [US/US]; 15013 S. 24th
Way, Phoenix, AZ 85048 (US).

(74) Agent: **ATKINS, Robert, D.**; Quarles & Brady Streich
Lang LLP, One Renaissance Square, Two North Central
Avenue, Phoenix, AZ 85004 (US).

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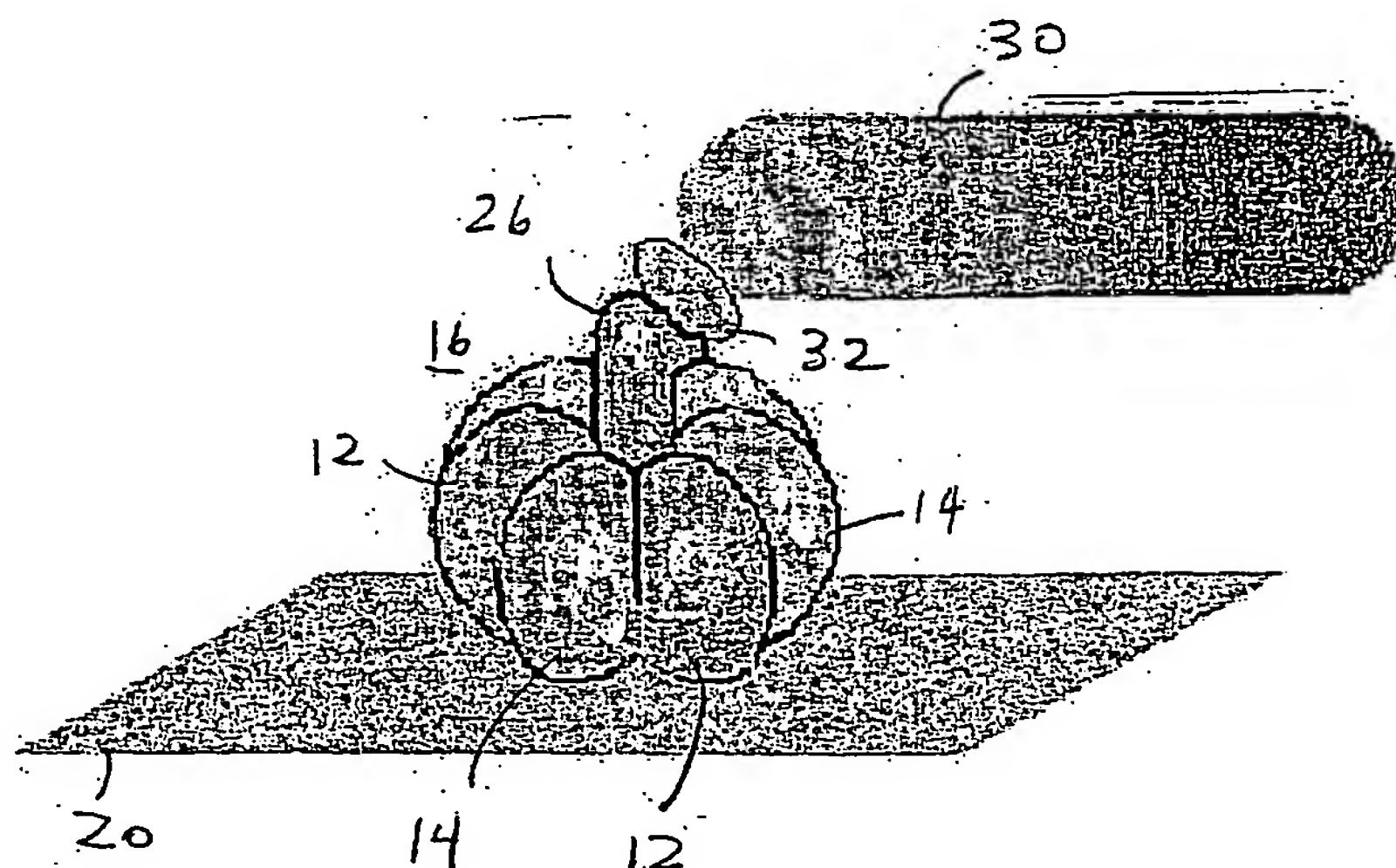
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(54) Title: **POLARIZATION-ENHANCED DETECTOR WITH GOLD NANORODS FOR DETECTING NANOSCALE ROTA-
TIONAL MOTION AND METHOD THEREFOR**



(57) Abstract: A nanoscale motion detector attaches a gold nanorod (30) to the rotating arm (26) of a molecular structure (10) to cause the nanoparticle to rotate. The molecular structure is an F1-ATPase enzyme. The gold nanorod is exposed to a light source. The long axis of the gold nanorod scatters red light when the nanorod is in a first position. The short axis of the gold nanorod scatters green light when the nanorod is in a second position. A polarizing filter filters the red and green light to detect the rotational motion by observing alternating red and green lights. A detection DNA stand (50) is coupled between the gold nanorod and the molecular structure. The detection DNA strand hybridizes with a target DNA strand (58) if the target DNA strand matches the detection DNA strand to form a structural link between the molecular structure and gold nanorod.

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**Polarization-Enhanced Detector with Gold Nanorods
For Detecting Nanoscale Rotational Motion and Method
Therefor**

5

Claim to Domestic Priority

[0001] The present non-provisional patent application claims priority to provisional application serial no. 60/432,589, entitled "Polarization-Enhanced Detector for
10 Nanoscale Rotational Motion using Gold Nanorods", filed on December 11, 2002.

Field of the Invention

15

[0002] The present invention relates in general to motion detectors and, more particularly, to a detector and method of detecting motion on the nanometer scale.

Background of the Invention

20

[0003] When working with man-made materials and naturally occurring substances, engineers and scientists are accustomed to observing and measuring physical characteristics and behavioral phenomena such as size,
25 shape, dimensions, transformation, motion, and other cause and effect relationships on the micrometer (μm) scale. For example, in the electronics and semiconductor industry, the dimensions of devices are measured in terms of microns. In biotechnology applications, most
30 instruments and techniques used to observe physical characteristics and behavioral phenomena, including microscopes, optical imaging, and electron micrographs are also in the micrometer range.

[0004] As technology advances, the dimensions of
35 devices, materials, and substances of interest continue

to dramatically shrink in size. Significant research and development is underway in what is commonly known as nanotechnology, i.e., devices and materials that exist and operate in the range of 1 to 1000 nanometers (nm).

5 It has been said that the ultimate refinement of realization and sensitivity is a single molecule. With nanotechnology, work is often done at the molecular level. Complex processes can take place in such a small space that the application become very portable.

10 Propagation times and energy consumption are negligible.

[0005] Nanotechnology finds applications in characterizing and monitoring nanoscopic systems ranging from single molecules to nano-electro-mechanical and nanofluidic systems. Researchers continue to look for
15 new applications of nanotechnology. The concept of realizing independently operating, man-made and engineered devices measured in terms of nanometers has become reality and will continue to progress. And, as the field of biotechnology advances, the need to observe,
20 measure, manipulate, control, and test substances and elements at the molecular level is ever more present.

[0006] One of the behavioral phenomena that exist in the world of nanotechnology is motion. Many aspects of the nanoworld are continuously in motion. The nature of
25 the motion is directly related to the physical characteristics and environment to which the nanoscale elements and structures are subjected. The ability to detect, observe, measure, and control such motion at the nanometer scale is important to the continuation of
30 research and development of new products and design methods. Modern instrumentation and research techniques have difficulty with the accurate and reliable detection of motion, particularly rotational motion, in the nanometer range.

[0007] Attempts have been made to detect and measure rotational motion of small particles by observing changes in the orientation of the particle over time under a microscope. For example, the F1-ATPase enzyme has been observed to exhibit rotational motion along its axis by using fluorescence microscopy to visualize time-dependent changes in the orientation of fluorescently labeled actin filaments, a protein which is about 0.5 to 4 μm long, attached to the rotating shaft of the enzyme. Other research has measured the rotation of anisotropically patterned fluorescent polymer microspheres in the range of 2 to 4 μm in diameter. The particles of interest possess sufficient anisotropy to allow its orientation to be seen under the microscope.

[0008] Alternatively, if a particle rotates about an axis that is not an axis of symmetry of the particle, then its rotation can be measured by tracking the centroid of the particle's image. The rotation of a 1 μm polystyrene sphere attached to the shaft of the F1-ATPase molecule has been measured by detecting the displacement of the centroid of the sphere's image due to the slight eccentricity in the shaft's rotation.

[0009] At the nanoscale, the direct approach to detecting rotational motion using a light-based microscope is usually ineffective, since the diffraction limit of the light makes it difficult to resolve features, and hence determine the orientation, of nanoscopic objects. That is, the magnitude of the change in position of the object is less than the diffraction limit of the light used to measure it.

[00010] The single-molecule fluorescence polarization spectroscopy and the centroid-tracking method have limitations in that the signal emitted by a single fluorophore is weak. Fluorescence polarization

spectroscopy requires a sensitive optical detection system. In addition, the probe is susceptible to photo-bleaching. Finally, the intensity of emission of a single fluorophore fluctuates. Single-molecule
5 fluorescence polarization spectroscopy cannot distinguish such fluctuations from those due to the rotation of the fluorophore, making the method susceptible to noise. The centroid-tracking method works only for off-axis rotation involves time-consuming, off-line image analysis. In
10 general, it is difficult to observe rotation of a circular object at any scale when viewed along the axis of rotation unless the rotation of the object is eccentric to the axis of rotation and/or the rotating object has an asymmetric shape.

15

Summary of the Invention

[00011] In one embodiment, the present invention is a method of detecting motion in nanoscale structures
20 comprising the steps of providing a molecular structure having a rotating arm, attaching a nanoparticle to the rotating arm of the molecular structure so that the nanoparticle rotates with the rotating arm of the molecular structure, exposing a light to the
25 nanoparticle, wherein a first surface of the nanoparticle scatters a first wavelength of the light when the nanoparticle is in a first position and a second surface of the nanoparticle scatters a second wavelength of the light when the nanoparticle is in a second position, and
30 filtering the first and second wavelengths of the light through a polarizing filter to detect rotational motion by observing alternating first and second wavelengths of the light.

[00012] In another embodiment, the present invention is

a method of detecting a substance comprising the steps of
attaching a detection DNA stand between a nanoparticle
and a rotating portion of a molecular structure,
hybridizing a target DNA strand corresponding to the
5 substance to be detected to the detection DNA strand if
the target DNA strand matches the detection DNA strand to
form a structural link between the nanoparticle and the
molecular structure, exposing a light to a first surface
of the nanoparticle to scatter a first wavelength of the
10 light, exposing a light to a second surface of the
nanoparticle to scatter a second wavelength of the light,
filtering the first and second wavelengths of the light,
and detecting presence of the substance upon observing
alternating first and second wavelengths of the filtered
15 light.

[00013] In yet another embodiment, the present
invention is a nanoscale motion detector comprising a
molecular structure having a rotating portion. A
nanoparticle is coupled to the rotating portion of the
20 molecular structure. A light source is incident to a
first surface of the nanoparticle to scatter a first
wavelength of the light when the nanoparticle is in a
first position and further incident to a second surface
of the nanoparticle to scatter a second wavelength of the
25 light when the nanoparticle is in a second position. A
polarizing filter filters the first and second
wavelengths of the light. The rotation motion is
detected by observing first and second wavelengths of the
filtered light.

Brief Description of the Drawings

[00014] FIG. 1 illustrates a representation of the F1-
ATPase enzyme;

FIG. 2 illustrates the various positions of the gold nanorod during one revolution of the gamma-subunit arm;

FIG. 3 is a dark field microscopy instrumentation setup used to detect the scattered light; and

FIG. 4 is another embodiment for detecting a target substance using rotation of the gold nanorods.

10 Detailed Description of the Drawings

[00015] The present invention relates to the detection and measurement of rotational motion at the nanometer level, that is, in the scale between 1 and 1000
15 nanometers, using anisotropic metal nanoparticle probes. The applications involved in characterizing and monitoring nanoscopic systems vary from single molecules to nano-electro-mechanical, nanofluidic systems, and other nanotechnologies. The present invention can be
20 used to observe the physical and behavioral characteristics of a single molecule. Another application provides for detection of the presence of specific substances.

[00016] Many enzymes are known to behave as molecular
25 motors, i.e., molecular structures that produce rotation motion. The molecular motors can be biological in origin or chemically synthesized. One such molecular motor is the F1-ATPase enzyme as represented in FIG. 1. F1-ATPase enzyme 10 includes alpha-subunits 12 and beta-subunits 14
30 in its base structure 16. Base structure 16 is non-rotating and anchored or attached preferentially to surface 20. Surface 20 can be part of a DNA microarray, glass slide, or other dielectric substrate. Gamma-subunit 26 is molecularly coupled to base structure 16

and oriented upward, normal to surface 20. F1-ATPase enzyme 10 operates as a molecular motor with the alpha-subunits 12 and beta-subunits 14 inducing rotation in gamma-subunit arm 26. Thus, gamma-subunit arm 26 behaves as a drive shaft, oriented upward, perpendicular to surface 20, and rotating in response to activity in alpha-subunits 12 and beta-subunits 14. The rotation of gamma-subunit arm 26 is difficult to directly see under a microscope.

[00017] A gold nanorod 30 is attached to gamma-subunit arm 26 with a protein or other bonding molecule 32, such as avidin. Gamma-subunit arm 26 can attach to any part of gold nanorod 30, e.g., at either end, in the middle, or any point in-between. The bonding molecule 32 links gold nanorod 30 to gamma-subunit arm 26 so that the rotational motion of gamma-subunit arm 26 is imparted to gold nanorod 30. Gold nanorod 30 spins around with the rotational motion of gamma-subunit arm 26.

[00018] Metal nanoparticles, such as gold nanorod 30, are efficient absorbers and scatterers of light owing to collective oscillations of their conduction electrons known as surface plasmons. The number, position, and shape of the surface plasmon bands are determined by the kind of metal, the size and shape of the particle, and the dielectric constant of the surrounding medium. Non-spherical nanoparticles possess multiple surface plasmon modes. For example, rod-shaped nanoparticles exhibit two resonant modes, corresponding to their long and short axes, respectively.

[00019] Gold nanorod 30 is shaped as a rod, shaft, or cylinder with rounded or flat ends. Gold nanorod 30 is about 15-30 nm in diameter and about 60-80 nm in length along its symmetrical axis. In other embodiments, gold nanorod 30 has a length-diameter aspect ratio between

2.5:1 and 20:1. The rod-shaped nanoparticle is typically grown from a base wafer or sphere. Gold nanorod 30 exhibits optical anisotropy in that it has two surface plasmon resonances that correspond to the diameter and length of the shaft. The short axis corresponds to the transverse plasmon resonance of the rod. The long axis corresponds to the longitudinal plasmon resonance of the rod. The short and long axes of gold nanorod 30 scatter incident white light at different wavelengths as a function of the respective dimensions. The longer axis has greater surface area than the shorter axis. The short axis or end surface of gold nanorod 30 scatters light at a shorter wavelength due to the lesser surface area and the long axis or side surface of gold nanorod 30 scatters light at a longer wavelength due to its larger surface area. In one embodiment, the short axis of gold nanorod 30 scatters green light having a wavelength of about 520-570 nm, while the long axis scatters red light having a wavelength of 685-730 nm. Other relative dimensions of gold nanorod 30 will scatter incident white light at two different wavelengths.

[00020] FIGs. 2a-2d illustrates a top view of gold nanorod 30 attached to F1-ATPase enzyme 10 at various positions A-D in one revolution of gamma-subunit arm 26. Drawing elements having a similar function are assigned the same reference numbers used in FIG. 1. In FIG. 2a, gold nanorod 30 is shown in position A with a first short axis or end surface oriented toward reference direction L. In FIG. 2b, a first long axis or side surface of gold nanorod 30 is shown in position B, oriented toward reference direction L. As the revolution of gamma-subunit 26 continues, a second short axis or end surface of gold nanorod 30 is shown in position C, oriented toward direction L, see FIG. 2c. In FIG. 2d, a second

long axis or side surface of gold nanorod 30 is shown in position D, oriented toward reference direction L.

[00021] The above-described optical scattering characteristics of gold nanorod 30, in combination with its rotational motion imparted by gamma-subunit arm 26, provides for observation and measurement of physical characteristics and behavioral phenomena of molecular structures on the nanoscale. In a simplified view, a full spectrum, white-light source is generally directed in reference direction L. The white light photons strike gold nanorod 30 as a wave function and, depending on the orientation of gold nanorod 30 with respect to the incident angle of the photons, some wavelengths of the light will be scattered. With the given dimensions of gold nanorod 30, when the nanorod is in position A, a green light is scattered. When nanorod 30 is in position B, a red light is scattered. When nanorod 30 is in position C, a green light is scattered. When nanorod 30 is in position D, a red light is scattered.

[00022] The scattered light from gold nanorod 30 is polarized. The intensities of the red and green resonances depend on the relative orientation of the nanorod to the plane of polarization of the incident light. The rotating nature of gold nanorod 30, due to its connection to the spinning F1-ATPase molecular motor, results in an observable light that flashes or blinks alternating green and red. The intensity and rate of blinking of the red and green light is a function of the speed of rotation of gamma-subunit arm 26. By observing the red and green blinking light from gold nanorod 30, the rotational motion of F1-ATPase enzyme 10 can be detected and measured. The visible blinking light is much easier to observe than the physical rotating structure itself. Moreover, the polarized nature of the

scattered wavelength causes gold nanorod 30 to go dark in between positions A-D. The flashing green and red lights are detectable, observable, and measurable using dark field microscopy.

5 **[00023]** A dark field microscopy instrumentation setup for detecting, observing, and measuring the scattered light from gold nanorod 30 is shown in FIG. 3. A full spectrum white light from light source 38 is incident to dark field condenser and lens 40, which in turn alters
10 the path of the light to create an oblique angle with respect to the long and short axes orientation of gold nanorod 30. The F1-ATPase enzyme 10 with attached gold nanorod 30 is positioned on slide surface 20. The F1-ATPase enzyme 10 and rotating gold nanorod 30 are exposed
15 to the white light. The rotational motion of the F1-ATPase enzyme 10, and the corresponding rotation of gold nanorod 30, cause the red and green wavelengths of the incident light to scatter depending on the orientation of the nanorod. Red light scatters when the long axis is
20 exposed to the white light and green light scatters when the short axis is exposed to the white light. The unscattered light continues on into objective lens 42 where it is blocked by iris 44. The wavelengths of the light, which are scattered by gold nanorod 30 pass
25 through iris or aperture 44.

30 **[00024]** A polarizing filter 46 is positioned at the output of objective lens 42 and passes the wavelength of the scattered light which is aligned with the polarizing filter, and further blocks any light, scattered or not, which is not aligned with the polarizing filter. The scattered red light, which is aligned with polarizing filter 46, passes through the filter. Likewise, the scattered green light, which is aligned with polarizing filter 46, passes through the filter. The intensity of

the red and green scattered light varies relative to the angle of polarization and has a maximum when the appropriate axis of gold nanorod 30 is parallel to the plane of polarization. When the light scattered from the long axis of gold nanorod 30 is aligned with polarizing filter 46 then red light is passed. When the light scattered from the short axis of gold nanorod 30 is aligned with polarizing filter 46 then green light is passed. No other light passes through polarizing filter 46, i.e., the output of polarizing filter 46 is otherwise dark.

[00025] The red and green lights are collected by optical processing equipment 48, which separates the red and green light into individual channels. MetaVue software is used to isolate, detect, observe, and measure the intensity of the red and green light. The F1-ATPase enzyme-induced rotation is observed as red and green blinking light through polarizing filter 46, due to surface plasmon resonance of gold nanorod 30. The alternating blinking red and green light has provided an observable and quantifiable representation of the physical and behavioral characteristics of the F1-ATPase enzyme 10, e.g., the blinking rate of the red and green light is a function of the speed of rotation of the F1-ATPase enzyme 10.

[00026] The gold nanorods can be used in a wide range of applications in nanoscience and nanotechnology where measurements of rotation are required. For example, the gold nanorod can be used to characterize and monitor the operation of biological nanomachines such as the F1-ATPase rotary biomolecular motor or man-made organic or nano-electro-mechanical devices. In addition, the gold nanorod is also suitable for measuring orientation, for example to sense whether a nanofluidic valve is in its

open or closed position. Other potential applications are in the area of microfluidics and nanofluidics, where the metal nanoparticles can be used as probes to measure the local velocity, vorticity, and shear rate of the flow field.

[00027] In using anisotropic metal nanoparticles, such as gold nanorods 30, to detect rotation, a variation in the intensity of the longitudinal plasmon resonance results when the excitation polarization is fixed and the particle is free to rotate. The intensity of the longitudinal resonance encodes the orientation of the nanoparticle. For example, a gold nanorod rotating in a homogeneous medium will appear alternately red and green when viewed under dark field linearly polarized white light illumination as a result of successive excitation of the longitudinal and transverse surface plasmon resonances, respectively. A nanorod rotating near a surface will also appear to blink red-green, since the transverse surface plasmon, being smaller in amplitude, will only be visible to the eye when the intensity of the longitudinal plasmon is low. Since the metal nanoparticles scatter light strongly, the variations in scattering intensity can be picked up with a conventional, non-intensified charge-coupled device (CCD) camera. Moreover, since the intensity of the transverse plasmon resonance is nearly constant, it serves as an internal reference against which changes in the longitudinal plasmon resonance can be measured.

[00028] Rotation is detected by measuring variations in the intensity of a diffraction-limited spot of light. The detection method is applicable to both on and off-axis rotation. The nanoparticle probe is not susceptible to photo-bleaching, nor does it exhibit the fluctuations characteristic of single fluorophore probes. Also, with

proper illumination, nanoparticles probes can be made to be much brighter than single fluorophores, which significantly reduces the demands on the sensitivity of the detection equipment or, alternatively, allows for the detection of much higher speed rotation. When the nanorod is rotating near a surface, the internal reference provided by the transverse plasmon resonance makes the method much more robust against sources of noise that cause variations in intensity of the longitudinal plasmon resonance unrelated to changes in orientation of the rod.

[00029] The nanoparticles can be ellipsoidal, rod-shaped, or other anisotropic shapes. The nanoparticles can be made of pure metals or alloys, and can be coated with a different type of metal or other material like glass. The nanoparticles can be flat structures patterned onto a metalized surface, for example by means of e-beam lithography.

[00030] In another embodiment, the gold nanorods are used as part of a molecular semaphore to detect the presence of a single molecule of a substance. Instead of attaching the gold nanorod directly to the gamma-subunit arm, a known deoxyribonucleic acid (DNA) oligonucleotide or an antibody is attached between gold nanorod 52 and the rotating gamma-subunit arm 26 of the F1-ATPase enzyme 10, as shown in FIG. 4. The known DNA structure is the detection DNA strand 50. A protein or bonding molecule 56, such as avidin, is used to anchor a first end of the detection DNA strand 50 to gamma-subunit arm 26 and further to anchor a second end of the detection DNA strand 50 to gold nanorod 52.

[00031] A sample of the target substance or molecule to be detected is introduced. The sequence of nucleotide base pairs is unique and defines the matching strand. If

the DNA sequence of the target molecule matches DNA strand 50, i.e., the nucleotide bases of the target DNA strand correspond to the nucleotide bases of the detection DNA strand 50, such as shown in FIG. 4, then the target DNA stand will hybridize to the detection DNA stand 50 to form a bridge or linkage. If the nucleotide bases of the target DNA strand do not correspond to the nucleotide bases of the detection DNA strand 50, then the DNA of the target molecule will not hybridize to the detection DNA strand 50 and no bridge or linkage is formed. The hybridized DNA structure is the bridge, which provides a structural link between gamma-subunit arm 26 and gold nanorod 52 to cause gold nanorod 52 to rotate with the F1-ATPase enzyme 10. Without the formation of the linkage, the gold nanorod 52 will not rotate with the F1-ATPase enzyme. Therefore, if the DNA strands match, i.e., if the target substance is detected, then the bridge is formed and gold nanorod 52 rotates with gamma-subunit arm 26. If the DNA strands do not match, i.e., the target substance is not detected, then the linkage is not formed and gold nanorod 52 does not rotate with gamma-subunit arm 26. If the target substance is detected, then the gold nanorod will rotate with the F1-ATPase enzyme and the nanoscale motion detector will flash alternating red and green lights as discussed above. If the target substance is not detected, then the gold nanorod will not rotate with the F1-ATPase enzyme and the detector will remain dark. The observation of the red and green lights is the detection mechanism for the target substance.

[00032] The present substance detection technique provides for detection using a single molecule of the target substance and can be used to detect specific proteins, diseases, and biological warfare agents like

Anthrax. Other applications include metabolites, proteomics, and a variety of drug testing.

[00033] A person skilled in the art will recognize that changes can be made in form and detail, and equivalents may be substituted for elements of the invention without departing from the scope and spirit of the invention. The present description is therefore considered in all respects to be illustrative and not restrictive, the scope of the invention being determined by the following claims and their equivalents as supported by the above disclosure and drawings.

Claims

What is claimed is:

- 5 1. A method of detecting motion in nanoscale structures, comprising:
 providing a molecular structure having a rotating arm;
 attaching a nanoparticle to the rotating arm of the
10 molecular structure so that the nanoparticle rotates with the rotating arm of the molecular structure;
 exposing a light to the nanoparticle, wherein a first surface of the nanoparticle scatters a first wavelength of the light when the nanoparticle is in a
15 first position and a second surface of the nanoparticle scatters a second wavelength of the light when the nanoparticle is in a second position; and
 filtering the first and second wavelengths of the light through a polarizing filter to detect rotational
20 motion by observing alternating first and second wavelengths of the light.
- 25 2. The method of claim 1, wherein the nanoparticle is rod-shaped.
- 30 3. The method of claim 2, wherein the nanoparticle is a gold nanorod.
4. The method of claim 1, wherein the first surface of the nanoparticle has greater area than the second surface of the nanoparticle.
5. The method of claim 4, wherein the first wavelength of the light is longer than the second wavelength of the

light.

5 6. The method of claim 5, wherein the first wavelength of the light is red light and the second wavelength of the light is green light.

7. The method of claim 1, wherein the molecular structure is an F1-ATPase enzyme.

10 8. The method of claim 1, further including the step of disposing a detection DNA stand between the nanoparticle and the molecular structure, wherein the detection DNA strand hybridizes with a target DNA strand, if the target DNA strand matches the detection DNA strand, to form a
15 structural link between the molecular structure and the nanoparticle.

9. A method of detecting motion in nanoscale structures, comprising:

20 attaching a nanoparticle to a rotating portion of a molecular structure;

exposing a light to a first surface of the nanoparticle to scatter a first wavelength of the light;

25 exposing a light to a second surface of the nanoparticle to scatter a second wavelength of the light; and

filtering the first and second wavelengths of the light to detect the rotational motion by observing the first and second wavelengths of the light.

30

10. The method of claim 9, wherein the nanoparticle is rod-shaped.

11. The method of claim 10, wherein the nanoparticle is

a gold nanorod.

5 12. The method of claim 9, wherein the first surface of the nanoparticle has greater area than the second surface of the nanoparticle.

10 13. The method of claim 12, wherein the first wavelength of the light is longer than the second wavelength of the light.

14. The method of claim 13, wherein the first wavelength of the light is red light and the second wavelength of the light is green light.

15 15. The method of claim 9, wherein the molecular structure is an F1-ATPase enzyme.

20 16. The method of claim 9, wherein the step of filtering the first and second wavelengths of the light uses a polarizing filter.

25 17. The method of claim 9, wherein the step of further including the step of disposing a detection DNA stand between the nanoparticle and the molecular structure, wherein the detection DNA strand hybridizes with a target DNA strand, if the target DNA strand matches the detection DNA strand, to form a structural link between the molecular structure and the nanoparticle.

30 18. A method of detecting a substance, comprising:
attaching a detection DNA stand between a nanoparticle and a rotating portion of a molecular structure;

hybridizing a target DNA strand corresponding to the

substance to be detected to the detection DNA strand if the target DNA strand matches the detection DNA strand to form a structural link between the nanoparticle and the molecular structure;

5 exposing a light to a first surface of the nanoparticle to scatter a first wavelength of the light; exposing a light to a second surface of the nanoparticle to scatter a second wavelength of the light; filtering the first and second wavelengths of the
10 light; and

detecting presence of the substance upon observing alternating first and second wavelengths of the filtered light.

15 19. The method of claim 18, wherein the nanoparticle is rod-shaped.

20. The method of claim 19, wherein the nanoparticle is a gold nanorod.

20

21. The method of claim 18, wherein the first surface of the nanoparticle has greater area than the second surface of the nanoparticle.

25 22. The method of claim 21, wherein the first wavelength of the light is longer than the second wavelength of the light.

30 23. The method of claim 22, wherein the first wavelength of the light is red light and the second wavelength of the light is green light.

24. The method of claim 18, wherein the step of filtering the first and second wavelengths of the light

uses a polarizing filter.

25. The method of claim 18, wherein the molecular structure is an F1-ATPase enzyme.

5

26. A method of detecting motion, comprising:
attaching an anisotropic particle to a rotating portion of a base structure;

10

exposing a light to the anisotropic particle to scatter first and second wavelengths of the light; and filtering the first and second wavelengths of the light to detect the rotation motion by observing the first and second wavelengths of the light.

15

27. The method of claim 26, wherein the anisotropic particle is rod-shaped.

28. The method of claim 27, wherein the anisotropic particle is a gold nanorod.

20

29. The method of claim 26, wherein the first surface of the anisotropic particle has greater area than the second surface of the anisotropic particle.

25

30. The method of claim 29, wherein the first wavelength of the light is longer than the second wavelength of the light.

30

31. The method of claim 30, wherein the first wavelength of the light is red light and the second wavelength of the light is green light.

32. The method of claim 26, wherein the base structure is an F1-ATPase enzyme.

33. A nanoscale motion detector, comprising:
a molecular structure having a rotating portion;
a nanoparticle coupled to the rotating portion of
5 the molecular structure;

a light source incident to a first surface of the
nanoparticle to scatter a first wavelength of the light
when the nanoparticle is in a first position and further
incident to a second surface of the nanoparticle to
10 scatter a second wavelength of the light when the
nanoparticle is in a second position; and

a polarizing filter filters the first and second
wavelengths of the light, wherein rotation motion is
detected by observing first and second wavelengths of the
15 filtered light.

34. The nanoscale motion detector of claim 33, wherein
the nanoparticle is rod-shaped.

20 35. The nanoscale motion detector of claim 34, wherein
the nanoparticle is a gold nanorod.

36. The nanoscale motion detector of claim 33, wherein
the molecular structure is an F1-ATPase enzyme.

25 37. The nanoscale motion detector of claim 33, further
including a detection DNA stand coupled between the
nanoparticle and the molecular structure, wherein the
detection DNA strand hybridizes with a target DNA strand,
30 if the target DNA strand matches the detection DNA
strand, to form a structural link between the molecular
structure and the nanoparticle.

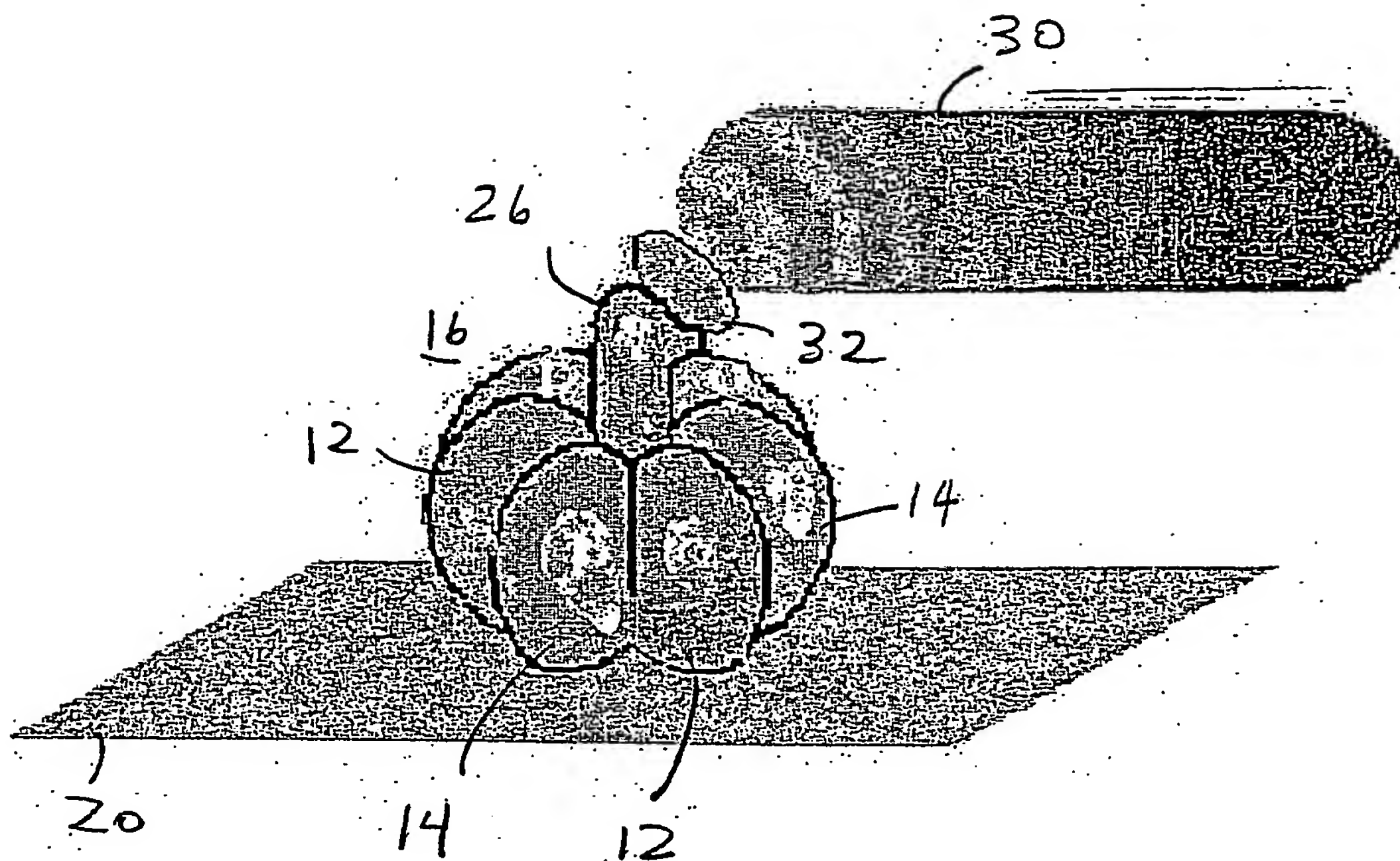


FIG. 1

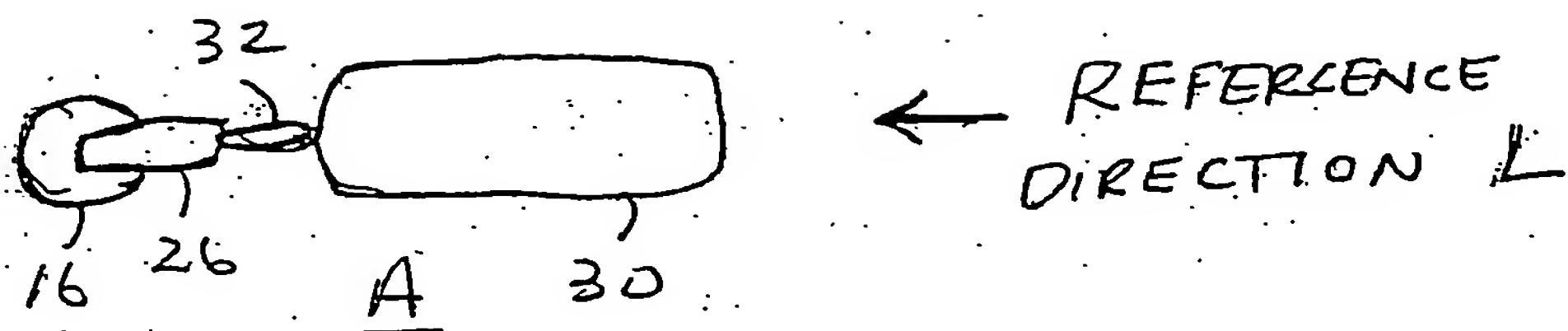


FIG. 2a

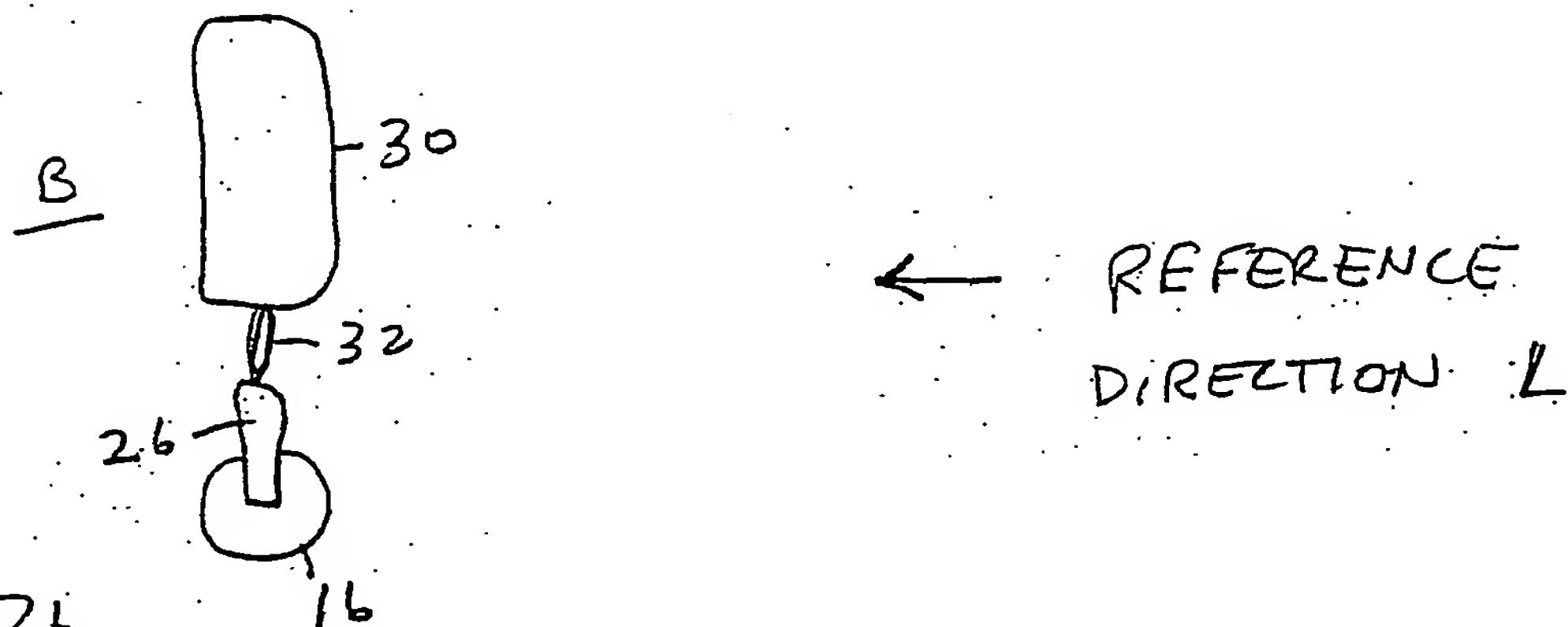


FIG. 2b

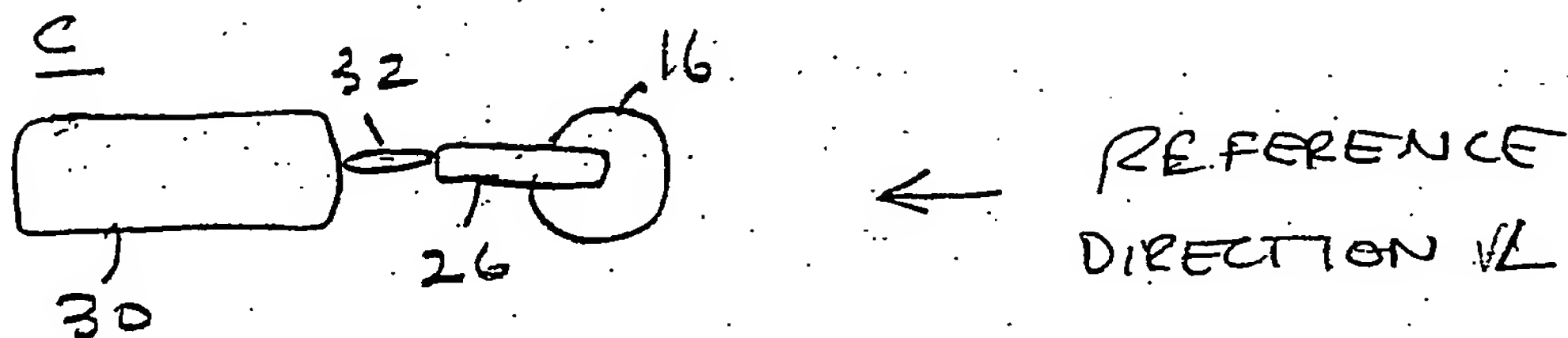


FIG. 3c

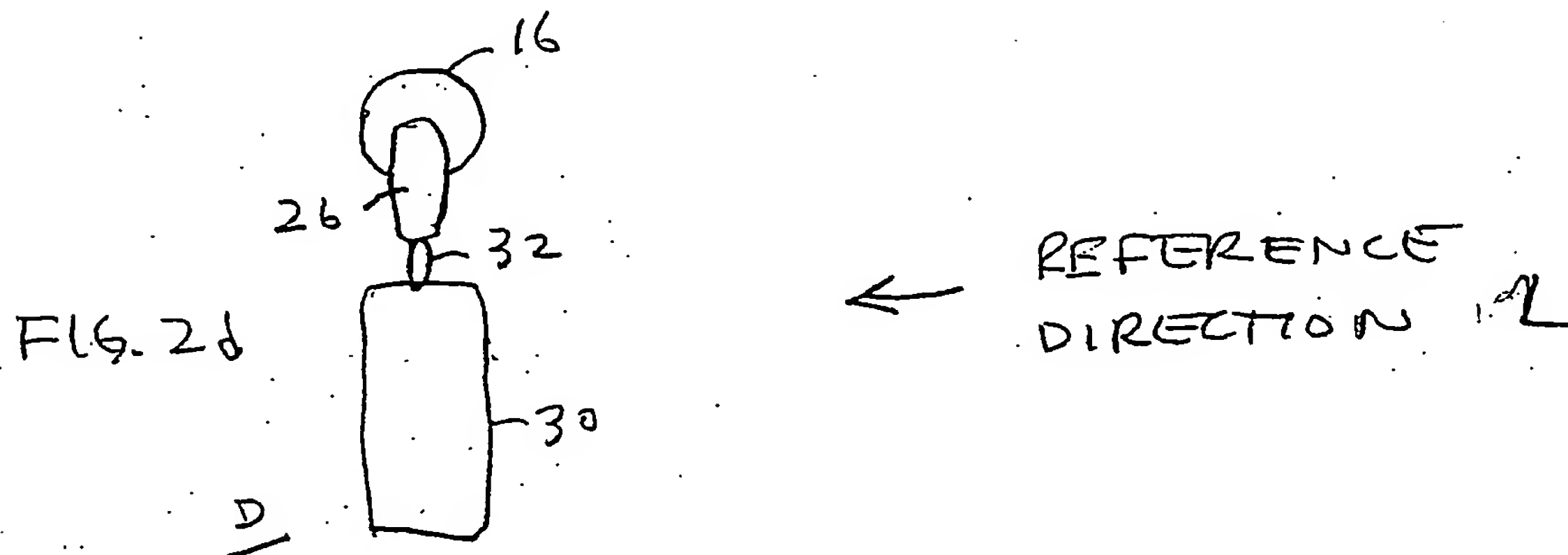


FIG. 2d

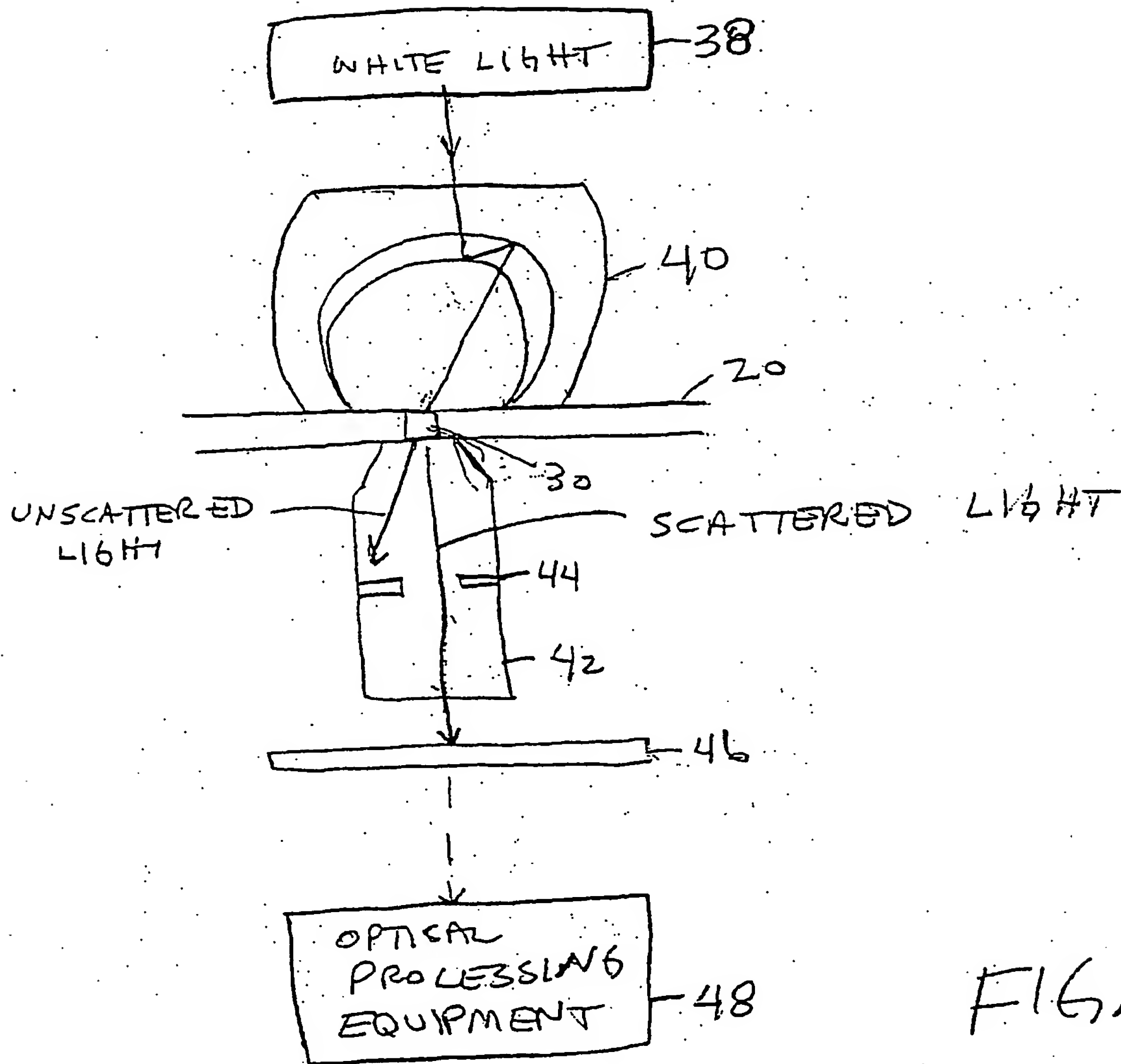


FIG. 3

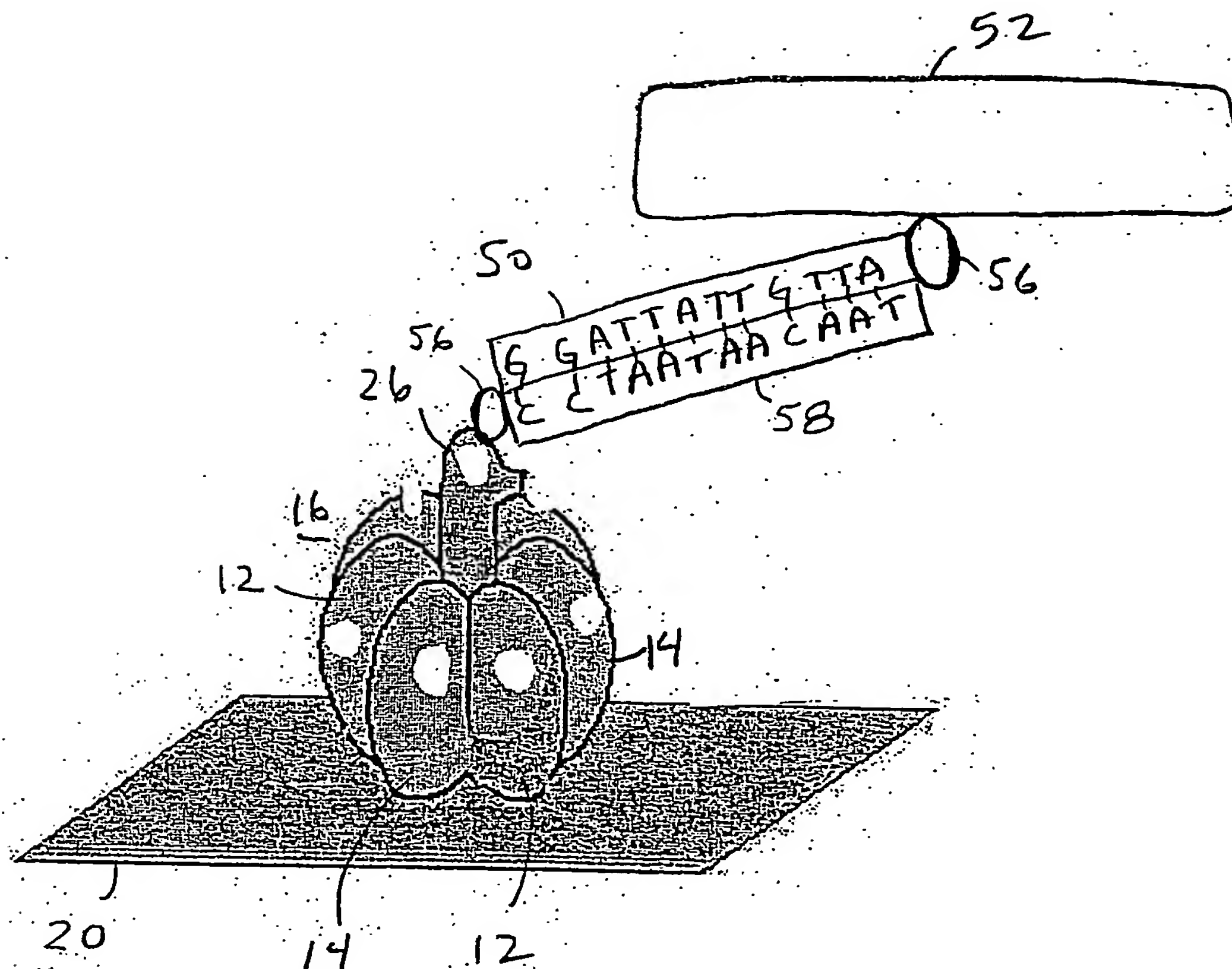


FIG. 4

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